Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the

application:

Listing of Claims:

1. (Original): A method of collecting a bodily fluid sample from an incision in the

skin comprising:

pressing against the skin a stimulator sleeve of a bodily fluid sampling device around the

incision to express the bodily fluid sample; and

moving a capillary tube of the bodily fluid sampling device towards the incision by

moving the capillary tube relative to the stimulator sleeve while the sleeve remains in contact

with the skin.

2. (Original): The method of claim 1, further comprising forming the incision in the

skin with a needle of the bodily fluid sampling device.

3. (Original): The method of claim 1, further comprising forming the incision with

the bodily fluid sampling device before said moving.

4. (Original): The method of claim 3, further comprising drawing the bodily fluid

from the incision into the capillary tube.

5. (Original): The method of claim 4, further comprising transferring the bodily fluid

onto a test strip located at one end of the capillary tube.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

Attorney Docket No. 7404-543; Document No. 309392

Page 2 of 22

6. (Original): The method of claim 5, further comprising analyzing the bodily fluid

on the test strip.

7. (Original): The method of claim 1, further comprising drawing the bodily fluid

from the incision into the capillary tube.

8. (Original): The method of claim 7, further comprising transferring the bodily fluid

from the capillary tube onto a test strip.

9. (Original): The method of claim 8, further comprising analyzing the bodily fluid

on the test strip.

10. (Original): A method of collecting a sample of bodily fluid from an incision in the

skin, comprising:

pressing against the skin a stimulator sleeve of a bodily fluid sampling device around the

incision to express at least a drop of the bodily fluid; and

moving a means for collecting the bodily fluid in the bodily fluid sampling device

towards the drop by moving the means for collecting the bodily fluid relative to the stimulator

sleeve while the sleeve remains in contact with the skin.

11. (Original): The method of claim 10, wherein:

the means for collecting the bodily fluid includes a capillary tube with an end; and

said moving includes extending the end of the capillary tube towards the drop.

12. (Original): The method of claim 10, wherein:

the bodily fluid sampling device includes an inner sleeve having a slot;

the stimulator sleeve is slidable relative to the inner sleeve;

the means for collecting the bodily fluid includes a test strip received in the slot of the

inner sleeve; and

said moving includes sliding the inner sleeve relative to the stimulator sleeve to contact

the test strip with the drop.

13. (Original): The method of claim 10, further comprising forming the incision with

the bodily fluid sampling device before said moving.

14. (Previously Presented): A method, comprising:

placing a sampling device in contact with a non-digit body part;

creating an incision in the non-digit body part with the sampling device; and

testing body fluid on the surface of the non-digit body part from the incision with the

sampling device while the sampling device remains in contact with the non-digit body part.

15. (Previously Presented): The method of claim 14, further comprising sampling the

body fluid from the incision with the sampling device before said testing.

16. (Previously Presented): The method of claim 15, wherein said sampling the body

fluid includes drawing fluid into a capillary in the sampling device via capillary action.

17. (Previously Presented): The method of claim 16, wherein said testing includes

analyzing the body fluid with a test strip disposed along the capillary.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

18. (Previously Presented): The method of claim 15, wherein said sampling includes:

moving a capillary from a first position where the capillary is displaced from the skin to a

second position where the capillary is adjacent the skin while the sampling device remains in

contact with the skin; and

drawing the body fluid from the incision into the capillary via capillary action.

19. (Previously Presented): The method of claim 14, further comprising said testing

includes analyzing the body fluid with a test strip disposed at an end of the sampling device

proximal the skin.

20. (Previously Presented): The method of claim 14, further comprising wherein the

non-digit body part is an earlobe or a limb.

21. (Previously Presented): A sampling module comprising:

a module body portion having a sampling site adjacent a lancet exit port where the

sharpened distal tip of the lancet exits a distal end of the module body portion that includes a

sample cavity in a distal end surface of the module body portion;

a lancet comprising a sharpened distal tip and shaft portion which is slidably disposed

within the module body portion and extendable from the lancet exit port; and

a sample reservoir in fluid communication with a sample cavity of the module body

portion.

22. (Previously Presented): The sampling module of claim 21 wherein a transverse

dimension of the sampling cavity is about 2 to about 5 times a transverse dimension of the lancet

shaft portion and wherein a sample flow channel is disposed between and in fluid

communication with the sample reservoir and the sample cavity.

23. (Previously Presented): The sampling module of claim 21 wherein the module

body portion is configured to be mechanically registered and secured adjacent a lancet driver.

24. (New): A tissue penetrating system, comprising:

a penetrating member driver;

a cartridge with a distal port and a proximal port and coupled to the penetrating member

driver;

an analyte detecting member coupled to a sample chamber, the analyte detecting member

being configured to determine a concentration of an analyte in a body fluid using a sample of a

body fluid disposed in the sample chamber;

a penetrating member with a sharpened distal tip and shaft portion that is slidably

disposed within the cartridge, wherein a tip of the penetrating member is configured to extend

through the opening of the sample chamber; and

a user interface configured to relay at least one of, skin penetrating performance or a skin

penetrating setting.

25. (New): The tissue penetrating system of claim 24, wherein the sample is less than

1 μ L of the body fluid.

26. (New): A tissue penetrating system, comprising:

a penetrating member driver;

a cartridge with a distal port and a proximal port and coupled to the penetrating member

driver;

an analyte detecting member coupled to a sample chamber, the analyte detecting member

being configured to determine a concentration of an analyte in a body fluid using a sample of a

body fluid disposed in the sample chamber;

a penetrating member with a sharpened distal tip and shaft portion that is slidably

disposed within the cartridge, wherein a tip of the penetrating member is configured to extend

through the opening of the sample chamber; and

a human interface providing at least one output.

27. (New): The tissue penetrating system of claim 26, wherein the sample is less than

 $1 \mu L$ of the body fluid.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

28. (New): The system of claim 26, wherein the at least one output is selected from, a

penetration event of a penetrating member, number of penetrating members remaining, time of

day, alarm, penetrating member trajectory waveform profile information, force for last

penetration event, the last penetration event, how or low battery status, analyte status, time to

change cassette status, jamming malfunction, and system status.

29. (New): The system of claim 26, wherein the human interface is selected from an

LED, an LED digital display, an LCD display, a sound generator, a buzzer, and a vibrating

device.

30. (New): The system of claim 26, wherein the housing is selected from at least one

of, a telephone, a watch, a PDA, electronic device, medical device, point of care device and a

decentralized diagnostic device.

31. (New): The system of claim 26, further comprising: an input device coupled to the

housing, the input device selected from one or more pushbuttons, a touch pad independent of the

display device, or a touch sensitive screen on a visual display.

32. (New): A skin penetrating system, comprising:

a housing member;

a penetrating member positioned in the housing member, and

an analyte detecting member coupled to a sample chamber, the analyte detecting member

being configured to determine a concentration of an analyte in a body fluid using a sample of a

body fluid disposed in the sample chamber, wherein a tip of the penetrating member is

configured to extend through an opening of the sample chamber.

33. (New): The skin penetrating system of claim 32, wherein the sample is less than 1

μL of the body fluid.

34. (New): The system of claim 32, further comprising:

a tissue stabilizer device coupled to the housing.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

35. (New): The system of claim 34, wherein the tissue stabilizer device is configured

to enhance fluid flow from a target tissue.

36. (New): The system of claim 34, wherein the tissue stabilizer device creates a

stretching of a skin surface.

37. (New): The system of claim 34, wherein the tissue stabilizer device is configured

to apply a force to a target tissue and cause the target tissue to press in an inward direction

relative to the housing member.

38. (New): The system of claim 34, wherein the tissue stabilizing member applies a

stimulation to a target tissue.

39. (New): The system of claim 32, wherein each penetrating member is an elongated

member without molded attachments.

40. (New): The system of claim 32, further comprising:

a support structure for receiving the penetrating members.

41. (New): A tissue penetrating system, comprising:

a penetrating member driver;

a cartridge with a distal port and a proximal port and coupled to the penetrating member

driver;

an analyte detecting member coupled to a sample chamber, the analyte detecting member

being configured to determine a concentration of an analyte in a body fluid using a sample of a

body fluid disposed in the sample chamber; and

a penetrating member with a sharpened distal tip and shaft portion that is slidably

disposed within the cartridge, wherein a tip of the penetrating member is configured to extend

through the opening of the sample chamber.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

42. (New): The tissue penetrating system of claim 41, wherein the sample is less than $1 \mu L$ of the body fluid.

43. (New): A method of penetrating a target tissue, comprising:

providing a tissue penetrating system with a penetrating member and an analyte detecting member coupled to a sample chamber,;

advancing a penetrating member through the target tissue; withdrawing the penetrating member from the target tissue. receiving a body fluid in the sample chamber.

44. (New): The method of claim 43, wherein no more than 1 μ L of the body fluid is received in the sample chamber.

45. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment; a coupler on said force generator configured to engage at least a portion of said elongate portion of the penetrating member and drive said member along a path into a tissue site and withdrawn from a tissue site;

an analyte detecting member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid; and

a user interface configured to relay at least one of, penetrating member performance or a penetrating member setting.

46. (New): A body fluid sampling system for use on a tissue site, the system

comprising:

Third Preliminary Amendment Application No. 10/612,852; Group Art Unit 3736 Attorney Docket No. 7404-543; Document No. 309392 Page 9 of 22

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator

moving said member along a path out of a housing having a penetrating member exit, into said

tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment; a

coupler on said force generator configured to engage at least a portion of said elongate portion of

the penetrating member and drive said member along a path into a tissue site and withdrawn

from a tissue site;

an analyte detecting member positioned to receive fluid from a wound created by said

penetrating member, said detection member configured to determine a concentration of an

analyte in the fluid using a sample of less than 1 mL of the fluid; and

a human interface providing at least one output.

47. (New): A body fluid sampling system for use on a tissue site, the system

comprising:

a drive force generator;;

a penetrating member operatively coupled to said force generator, said force generator

moving said member along a path out of a housing having said a penetrating member exit, into

said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

a skin stabilizer device suitable for stretching a surface of a tissue site, said skin stabilizer

at least partially surrounding the penetrating member exit;

an analyte detecting member positioned to receive fluid from a wound created by said

penetrating member, said detection member configured to determine a concentration of an

analyte in the fluid using a sample of less than 1 mL of the fluid; and

a user interface configured to relay at least one of, penetrating member performance or a

penetrating member setting.

48. (New): A body fluid sampling system for use on a tissue site, the system

comprising:

a drive force generator;;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having said a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

a skin stabilizer device suitable for stretching a surface of a tissue site, said skin stabilizer at least partially surrounding the penetrating member exit;

an analyte detecting member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid; and

a human interface providing at least one output.

49. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having said a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

an analyte detecting member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid; and

a user interface configured to relay at least one of, penetrating member performance or a penetrating member setting..

50. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having said a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

an analyte detecting member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid; and

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736 Attorney Docket No. 7404-543; Document No. 309392

Page 11 of 22

a human interface providing at least one output.

51. (New): A body fluid sampling system for use on a tissue site, the system

comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator

moving said member along a path out of a housing having a penetrating member exit, into said

tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment.

a coupler on said force generator configured to engage at least a portion of said elongate

portion of the penetrating member and drive said member along a path into a tissue site and

withdrawn from a tissue site;

a skin stabilizing member associated with said housing and positioned to at least partially

surround an impact location of the penetrating member on the tissue site;

a user interface configured to relay at least one of, penetrating member performance or a

penetrating member setting.

52. (New): A body fluid sampling system for use on a tissue site, the system

comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator

moving said member along a path out of a housing having a penetrating member exit, into said

tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment.

a coupler on said force generator configured to engage at least a portion of said elongate

portion of the penetrating member and drive said member along a path into a tissue site and

withdrawn from a tissue site;

a skin stabilizing member associated with said housing and positioned to at least partially

surround an impact location of the penetrating member on the tissue site; and

a human interface providing at least one output.

53. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having said a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

a skin stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site;

a user interface configured to relay at least one of, penetrating member performance or a penetrating member setting.

54. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having said a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

a skin stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site; and

a human interface providing at least one output.

55. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

an analyte detection member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid;

a skin stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site;

a user interface configured to relay at least one of, penetrating member performance or a penetrating member setting.

56. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

an analyte detection member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid;

a skin stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site; and

a human interface providing at least one output.

57. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment; a coupler on said force generator configured to engage at least a portion of said elongate portion of the penetrating member and drive said member along a path into a tissue site and withdrawn from a tissue site;

an analyte detecting member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid.

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736 Attorney Docket No. 7404-543; Document No. 309392

Page 14 of 22

58. (New): A method of obtaining a sample of capillary whole blood from a target tissue, comprising

providing a penetrating system that includes a tissue stabilizing member; applying skin stimulation to a skin surface site with the tissue stabilizing member; introducing a penetrating member through the skin surface site to form an incision; and collecting blood from the incision in the penetrating system.

- 59. (New): The method of claim 58, wherein the skin stimulation increases blood circulation at the skin surface.
 - 60. (New): A tissue penetrating device, comprising: a housing;

at least one penetrating member a penetrating member driver coupled to the at least one penetrating member;

a tissue stabilizer member coupled to the housing; and a human interface providing at least one output.

- 61. (New): The system of claim 60, wherein the at least one output is selected from, a penetration event of a penetrating member, number of penetrating members remaining, time of day, alarm, penetrating member trajectory waveform profile information, force for last penetration event, the last penetration event, how or low battery status, analyte status, time to change cassette status, jamming malfunction, and system status.
- 62. (New): The system of claim 60, wherein the human interface is selected from an LED, an LED digital display, an LCD display, a sound generator, a buzzer, and a vibrating device.
- 63. (New): The system of claim 60, wherein the housing is selected from at least one of, a telephone, a watch, a PDA, electronic device, medical device, point of care device and a decentralized diagnostic device.

Third Preliminary Amendment Application No. 10/612,852; Group Art Unit 3736 Attorney Docket No. 7404-543; Document No. 309392 Page 15 of 22 64. (New): The system of claim 60, further comprising:

an input device coupled to the housing, the input device selected from one or more pushbuttons, a touch pad independent of the display device, or a touch sensitive screen on a visual display.

- 65. (New): The system of claim 60, further comprising:
- a data exchange device for coupling the tissue penetrating system to support equipment.
- 66. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

wherein said penetrating member is an elongate member without a molded attachment.

a coupler on said force generator configured to engage at least a portion of said elongate portion of the penetrating member and drive said member along a path into a tissue site and withdrawn from a tissue site;

a tissue stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site

67. (New): A body fluid sampling system for use on a tissue site, the system comprising:

a drive force generator;

a penetrating member operatively coupled to said force generator, said force generator moving said member along a path out of a housing having a penetrating member exit, into said tissue site, stopping in said tissue site, and withdrawing out of said tissue site;

an analyte detection member positioned to receive fluid from a wound created by said penetrating member, said detection member configured to determine a concentration of an analyte in the fluid using a sample of less than 1 mL of the fluid;

Third Preliminary Amendment

Application No. 10/612,852; Group Art Unit 3736

Attorney Docket No. 7404-543; Document No. 309392

Page 16 of 22

a tissue stabilizing member associated with said housing and positioned to at least partially surround an impact location of the penetrating member on the tissue site.